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110TH CONGRESS
1ST SESSION

H. R. 962

To amend the Federal Food, Drug, and Cosmetic Act to preserve the effectiveness of medically important antibiotics used in the treatment of human and animal diseases.

IN THE HOUSE OF REPRESENTATIVES

FEBRUARY 8, 2007

Ms. SLAUGHTER introduced the following bill; which was referred to the Committee on Energy and Commerce

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A BILL

To amend the Federal Food, Drug, and Cosmetic Act to preserve the effectiveness of medically important antibiotics used in the treatment of human and animal diseases.

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

SECTION 1. SHORT TITLE; TABLE OF CONTENTS.

(a) SHORT TITLE.—This Act may be cited as the ''Preservation of Antibiotics for Medical Treatment Act of 2007''.

(b) TABLE OF CONTENTS.—The table of contents of this Act is as follows:

SEC. 2. FINDINGS.

The Congress finds that—

(1)(A) in January 2001, a Federal interagency task force released an action plan to address the continuing decline in effectiveness of antibiotics against common bacterial infections, referred to as antibiotic resistance;

(B) the task force determined that antibiotic resistance is a growing menace to all people and poses a serious threat to public health; and

(C) the task force cautioned that if current trends continue, treatments for common infections will become increasingly limited and expensive, and, in some cases, nonexistent;

(2) antibiotic resistance, resulting in a reduced number of effective antibiotics, may significantly impair the ability of the United States to respond to terrorist attacks involving bacterial infections or a large influx of hospitalized patients;

(3)(A) any overuse or misuse of antibiotics contributes to the spread of antibiotic resistance, whether in human medicine or in agriculture; and

(B) recognizing the public health threat caused by antibiotic resistance, Congress took several steps to curb antibiotic overuse in human medicine through amendments to the Public Health Service Act (42 U.S.C. 201 et seq.) made by section 102 of the Public Health Threats and Emergencies Act (Public Law 106–505, title I; 114 Stat. 2315), but has not yet addressed antibiotic overuse in agriculture;

(4) in a March 2003 report, the National Academy of Sciences stated that—

(A) a decrease in antimicrobial use in human medicine alone will have little effect on the current situation; and

(B) substantial efforts must be made to decrease inappropriate overuse in animals and agriculture;

(5)(A) an estimated 70 percent of the antibiotics and other antimicrobial drugs used in the United States are fed to farm animals for nontherapeutic purposes, including—

(i) growth promotion; and

(ii) compensation for crowded, unsanitary, and stressful farming and transportation conditions; and

(B) unlike human use of antibiotics, these nontherapeutic uses in animals typically do not require a prescription;

(6)(A) many scientific studies confirm that the nontherapeutic use of antibiotics in agricultural animals contributes to the development of antibiotic-resistant bacterial infections in people;

(B) the periodical entitled ''Clinical Infectious Diseases'' published a report in June 2002, based on a 2-year review by experts in human and veterinary medicine, public health, microbiology, biostatistics, and risk analysis, of more than 500 scientific studies on the human health impacts of antimicrobial use in agriculture; and

(C) the report recommended that antimicrobial agents should no longer be used in agriculture in the absence of disease, but should be limited to therapy for diseased individual animals and prophylaxis when disease is documented in a herd or flock;

(7) the United States Geological Survey reported in March 2002 that—

(A) antibiotics were present in 48 percent of the streams tested nationwide; and

(B) almost half of the tested streams were downstream from agricultural operations;

(8) an April 1999 study by the General Accounting Office concluded that resistant strains of 3 microorganisms that cause food-borne illness or disease in humans—Salmonella, Campylobacter, and E. coli—are linked to the use of antibiotics in animals;

(9)(A) in January 2003, Consumer Reports published test results on poultry products bought in grocery stores nationwide showing disturbingly high levels of Campylobacter and Salmonella bacteria that were resistant to antibiotics used to treat food-borne illnesses; and

(B) further studies showed similar results in other meat products;

(10) in October 2001, the New England Journal of Medicine published an editorial urging a ban on nontherapeutic use of medically important antibiotics in animals;

(11)(A) in 1999, the European Union banned the practice of feeding medically important antibiotics to animals other than for disease treatment or control, and prior to that, individual European

countries had banned the use of specific antibiotics in animal feed; and

(B) those countries have experienced no significant impact on animal health or productivity, food safety, or meat prices, and more importantly, levels of resistant bacteria have declined sharply;

(12) in 1998, the National Academy of Sciences noted that antibiotic-resistant bacteria generate a minimum of $4,000,000,000 to $5,000,000,000 in costs to United States society and individuals yearly;

(13) a year later, the National Academy of Sciences estimated that eliminating the use of all antibiotics as feed additives would cost each American consumer less than $5 to $10 per year;

(14) the American Medical Association, the American Public Health Association, the National Association of County and City Health Officials, and the National Campaign for Sustainable Agriculture, are among the more than 300 organizations representing health, consumer, agricultural, environmental, humane, and other interests that support enactment of legislation to phase out nontherapeutic use in farm animals of medically important antibiotics;

(15) the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.)—

(A) requires that all drugs be shown to be safe before the drugs are approved; and

(B) places the burden on manufacturers to account for health consequences and prove safety;

(16)(A) the Food and Drug Administration recently modified the drug approval process for antibiotics to recognize the development of resistant bacteria as an important aspect of safety;

(B) however, most antibiotics currently used in animal production systems for nontherapeutic purposes were approved before the Food and Drug Administration began giving in-depth consideration to resistance during the drug-approval process; and

(C) the Food and Drug Administration has not established a schedule for reviewing those existing approvals; and

(17) certain non-routine uses of antibiotics in animal agriculture are legitimate to prevent animal disease; and

(18)(A) an April 2004 study by the General Accounting Office concluded that Federal agencies do not collect the critical data on antibiotic use in ani-

mals that they need to support research on human health risks; and

(B) the report recommends that the Department of Agrigulture and the Department of Health and Human Services develop and implement a plan to collect data on antibiotic use in animals.

SEC. 3. PURPOSE.

The purpose of this Act is to preserve the effectiveness of medically important antibiotics used in the treatment of human and animal diseases by phasing out use of certain antibiotics for nontherapeutic purposes in food-producing animals.

TITLE I—SAFETY OF CRITICAL ANTIMICROBIAL ANIMAL DRUGS

SEC. 101. PROOF OF SAFETY OF CRITICAL ANTIMICROBIAL ANIMAL DRUGS.

(a) DEFINITIONS.—Section 201 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321) is amended by adding at the end the following:

''(rr) CRITICAL ANTIMICROBIAL ANIMAL DRUG.—The term 'critical antimicrobial animal drug' means a drug that—

''(1) is intended for use in food-producing animals; and

''(2) is composed wholly or partly of—

''(A) any kind of penicillin, tetracycline, macrolide, lincosamide, streptogramin, aminoglycoside, or sulfonamide; or

''(B) any other drug or derivative of a drug that is used in humans or intended for use in humans to treat or prevent disease or infection caused by microorganisms.

''(ss) NONTHERAPEUTIC USE.—The term 'nontherapeutic use', with respect to a critical antimicrobial animal drug, means any use of the drug as a feed or water additive for an animal in the absence of any clinical sign of disease in the animal for growth promotion, feed efficiency, weight gain, routine disease prevention, or other routine purpose.''.

(b) APPLICATIONS PENDING OR SUBMITTED AFTER ENACTMENT.—Section 512(d)(1) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(d)(1)) is amended—

(1) in the first sentence—

(A) in subparagraph (H), by striking ''or'' at the end;

(B) by redesignating subparagraph (I) as subparagraph (J); and

(C) by inserting after subparagraph (H) the following:

''(I) with respect to a critical antimicrobial animal drug or a drug of the same chemical class as a critical antimicrobial animal drug, the applicant has failed to demonstrate that there is a reasonable certainty of no harm to human health due to the development of antimicrobial resistance that is attributable, in whole or in part, to the nontherapeutic use of the drug; or''; and

(2) in the second sentence, by striking ''(A) through (I)'' and inserting ''(A) through (J)''.

(c) PHASED ELIMINATION OF NONTHERAPEUTIC USE IN ANIMALS OF CRITICAL ANTIMICROBIAL ANIMAL DRUGS IMPORTANT FOR HUMAN HEALTH.—Section 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b) is amended by adding at the end the following:

''(q) PHASED ELIMINATION OF NONTHERAPEUTIC USE IN ANIMALS OF CRITICAL ANTIMICROBIAL ANIMAL DRUGS IMPORTANT FOR HUMAN HEALTH.—

''(1) APPLICABILITY.—This subsection applies to the nontherapeutic use in a food-producing animal of a drug—

''(A)(i) that is a critical antimicrobial animal drug; or

"(ii) that is of the same chemical class as a critical antimicrobial animal drug; and

"(B)(i) for which there is in effect an approval of an application or an exemption under subsection (b), (i), or (j) of section 505; or

"(ii) that is otherwise marketed for use.

"(2) WITHDRAWAL.—The Secretary shall withdraw the approval of a nontherapeutic use in food-producing animals described in paragraph (1) on the date that is 2 years after the date of enactment of this subsection unless—

"(A) before the date that is 2 years after the date of the enactment of this subsection, the Secretary makes a final written determination that the holder of the approved application has demonstrated that there is a reasonable certainty of no harm to human health due to the development of antimicrobial resistance that is attributable in whole or in part to the nontherapeutic use of the drug; or

"(B) before the date specified in subparagraph (A), the Secretary makes a final written determination under this subsection, with respect to a risk analysis of the drug conducted by the Secretary and other relevant informa-

tion, that there is a reasonable certainty of no harm to human health due to the development of antimicrobial resistance that is attributable in whole or in part to the nontherapeutic use of the drug.

''(3) EXEMPTIONS.—Except as provided in paragraph (5), if the Secretary grants an exemption under section 505(i) for a drug that is a critical antimicrobial animal drug, the Secretary shall rescind each approval of a nontherapeutic use in a food-producing animal of the critical antimicrobial animal drug, or of a drug in the same chemical class as the critical antimicrobial animal drug, as of the date that is 2 years after the date on which the Secretary grants the exemption.

''(4) APPROVALS.—Except as provided in paragraph (5), if an application for a drug that is a critical antimicrobial animal drug is submitted to the Secretary under section 505(b), the Secretary shall rescind each approval of a nontherapeutic use in a food-producing animal of the critical antimicrobial animal drug, or of a drug in the same chemical class as the critical antimicrobial animal drug, as of the date that is 2 years after the date on which the application is submitted to the Secretary.

''(5) EXCEPTION.—Paragraph (3) or (4), as the case may be, shall not apply if—

''(A) before the date on which approval would be rescinded under that paragraph, the Secretary makes a final written determination that the holder of the application for the approved nontherapeutic use has demonstrated that there is a reasonable certainty of no harm to human health due to the development of antimicrobial resistance that is attributable in whole or in part to the nontherapeutic use in the food-producing animal of the critical antimicrobial animal drug; or

''(B) before the date specified in subparagraph (A), the Secretary makes a final written determination under this subsection, with respect to a risk analysis of the critical antimicrobial animal drug conducted by the Secretary and any other relevant information, that there is a reasonable certainty of no harm to human health due to the development of antimicrobial resistance that is attributable in whole or in part to the nontherapeutic use of the drug.''.

TITLE II—USE OF CRITICAL ANTIMICROBIAL ANIMAL DRUGS IN AGRICULTURE

SEC. 201. COLLECTION OF DATA ON CRITICAL ANTIMICROBIAL ANIMAL DRUGS.

(a) IN GENERAL.—Chapter V of the Federal Food, Drug, and Cosmetic Act is amended by inserting after section 512 (21 U.S.C. 360b) the following:

"SEC. 512A. COLLECTION OF DATA ON CRITICAL ANTIMICROBIAL ANIMAL DRUGS.

''(a) IN GENERAL.—Not later than July 1 of each year, a manufacturer of a critical antimicrobial animal drug or an animal feed for food-producing animals bearing or containing a critical antimicrobial animal drug shall submit to the Secretary a report, in such form as the Secretary shall require, containing information on the sales during the previous calendar year of the critical antimicrobial animal drug or the animal feed.

''(b) INFORMATION TO BE INCLUDED.—A report under subsection (a) shall—

''(1) state separately the quantity of the critical antimicrobial animal drug, including such quantity in animal feed bearing or containing the critical antimicrobial drug, sold for each kind of food-producing animal;

''(2) describe the claimed purpose of use for the drug for each kind of food-producing animal as being for growth promotion, weight gain, feed efficiency, disease prevention, disease control, disease treatment, or another purpose; and

''(3) describe the dosage form of the drug.

''(c) PUBLICATION.—

''(1) IN GENERAL.—The Secretary shall make the information submitted under subsection (a) available to the public not less than annually.

''(2) PROTECTION OF CONFIDENTIALITY.—The Secretary may aggregate information, if necessary, so as to avoid disclosure under paragraph (1) of confidential business information.''.

(b) VIOLATION.—Subsection (e) of section 301 of the Federal Food, Drug and Cosmetic Act (21 U.S.C. 331(e)) is amended by striking ''515(f)'' and inserting ''512A, 515(f)''.

(c) EFFECTIVE DATE.—The amendments made by this section shall take effect on January 1, 2008.

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